EC Certificate

Directive 93/42/EEC Annex II, excluding Section 4
Full Quality Assurance System
Medical Devices

Registration No.: HD 60131226 0001

Report No.: 15078953 004

Manufacturer:
Jiangsu Ate Medical Technology
Co., Ltd.
No. 8, Lanxiang Road
Wujin Economic Development Zone
213161 Jiangsu
China

Products:
Disposable Injection Needles, Disposable Grasping Forceps,
Disposable Biopsy Forceps, Disposable Spray Pipes,
Disposable Hemostatic Clips, Disposable Polypectomy Snares

Replaces Approval, Registration No.: HD 60109072 0001

Expiry Date: 2023-04-13

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date: 2018-07-27

Date: 2018-07-27

Notified Body

X. Ren

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

Digitally signed by Grabazei Alexandru
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